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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/580,786	08/22/2007	Goran Nilsson	D7873.0002	7122
32172 DICKSTEIN SI	7590 04/22/201 HAPIRO LLP	EXAMINER		
1633 Broadway	,	CHICKOS, SARAH J		
NEW YORK, NY 10019			ART UNIT	PAPER NUMBER
			1619	
			MAIL DATE	DELIVERY MODE
			04/22/2011	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)
	10/580,786	NILSSON ET AL.
Office Action Summary	Examiner	Art Unit
	SARAH CHICKOS	1619
The MAILING DATE of this communication app Period for Reply	pears on the cover sheet with t	he correspondence address
A SHORTENED STATUTORY PERIOD FOR REPL WHICHEVER IS LONGER, FROM THE MAILING D - Extensions of time may be available under the provisions of 37 CFR 1.1 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period - Failure to reply within the set or extended period for reply will, by statute Any reply received by the Office later than three months after the mailin earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICAT 136(a). In no event, however, may a reply but will apply and will expire SIX (6) MONTHS a, cause the application to become ABAND	TON. De timely filed from the mailing date of this communication. ONED (35 U.S.C. § 133).
Status		
 Responsive to communication(s) filed on <u>25 M</u> This action is FINAL. Since this application is in condition for alloward closed in accordance with the practice under M 	s action is non-final. nce except for formal matters,	·
Disposition of Claims		
4) ☑ Claim(s) 1-16 and 18-21 is/are pending in the 4a) Of the above claim(s) is/are withdra 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) is/are rejected. 7) ☐ Claim(s) is/are objected to. 8) ☑ Claim(s) 1-16 and 18-21 are subject to restrict	wn from consideration.	nt.
Application Papers		
9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) accomplicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Examine 11.	cepted or b) objected to by t drawing(s) be held in abeyance. tion is required if the drawing(s) is	See 37 CFR 1.85(a). s objected to. See 37 CFR 1.121(d).
Priority under 35 U.S.C. § 119		
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the priority application from the International Burea * See the attached detailed Office action for a list	ts have been received. ts have been received in Appli prity documents have been rec u (PCT Rule 17.2(a)).	cation No eived in this National Stage
Attachment(s)		
1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date	4) Interview Sumn Paper No(s)/Ma 5) Notice of Inform 6) Other:	

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DETAILED ACTION

Election/Restrictions

Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group I, claims 2-7, 10-16 and 18-21 drawn to a pharmaceutical composition for oral administration comprising isotretinoin, a lipid material comprising membrane lipids and non-polar lipids.

Group II, claims 2-6, 8, 10-16 and 18-21 drawn to a pharmaceutical composition for oral administration comprising an immunosuppressant, a lipid material comprising membrane lipids and monoglycerides.

Group III, claims 2-6, 9-16 and 18-21 drawn to a pharmaceutical composition for oral administration comprising an antiviral, a lipid material comprising membrane lipids, and monoglycerides.

The inventions listed as Groups I-III do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: They are not joined by a special technical feature.

The technical feature that is shared by the groups is the pharmaceutical composition for oral administration comprising an active substance having a food effect in combination with a lipid material comprising membrane lipids. This composition is already known in the art as disclosed by Herslof et al. WO 00/32219, published June 8, 2000. (IDS 5/25/2006). WO00/32219 discloses a pharmaceutical composition comprising a cyclosporin as an active substance in a lipid carrier wherein the lipid carrier comprises membrane lipid in combination with monoglycerides and non-polar lipids. The composition contains 0.5-25% cyclosporine, 10-45% membrane lipids, 10-55% monoglycerides, 0-45% non-polar lipids (See page 3).

The expression "special technical feature" refers to those features that define a contribution which each of the claimed inventions, considered as a whole, makes over

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the prior art. Thus, a feature found in the prior art cannot be considered to be a special technical feature.

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1. Claim 1 link(s) inventions I, II and III. The restriction requirement separating the linked inventions is **subject to** the nonallowance of the linking claim(s), claim 1. Upon the indication of allowability of the linking claim(s), the restriction requirement as to the linked inventions **shall** be withdrawn and any claim(s) depending from or otherwise requiring all the limitations of the allowable linking claim(s) will be rejoined and fully examined for patentability in accordance with 37 CFR 1.104. **Claims that require all the limitations of an allowable linking claim** will be entered as a matter of right if the amendment is presented prior to final rejection or allowance, whichever is earlier. Amendments submitted after final rejection are governed by 37 CFR 1.116; amendments submitted after allowance are governed by 37 CFR 1.312.

Applicant(s) are advised that if any claim presented in a continuation or divisional application is anticipated by, or includes all the limitations of, the allowable linking claim, such claim may be subject to provisional statutory and/or nonstatutory double patenting rejections over the claims of the instant application. Where a restriction requirement is withdrawn, the provisions of 35 U.S.C. 121 are no longer applicable. *In re Ziegler*, 443 F.2d 1211, 1215, 170 USPQ 129, 131-32 (CCPA 1971). See also MPEP § 804.01.

Applicant is advised that the reply to this requirement to be complete must include (i) an election of a invention to be examined even though the requirement may be traversed (37 CFR 1.143) and (ii) identification of the claims encompassing the elected invention.

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The election of an invention may be made with or without traverse. To reserve a right to petition, the election must be made with traverse. If the reply does not distinctly and specifically point out supposed errors in the restriction requirement, the election shall be treated as an election without traverse. Traversal must be presented at the time of election in order to be considered timely. Failure to timely traverse the requirement will result in the loss of right to petition under 37 CFR 1.144. If claims are added after the election, applicant must indicate which of these claims are readable on the elected invention.

Election of Species Applicable to Groups II and III Above

2. If Group II or III is elected, the following species elections are applicable.

Claim 8 is generic to the following disclosed patentably distinct species: the immunosuppressants that are disclosed but not claimed are listed on page 10 beginning at line 40 or elsewhere in the specification. Claim 9 is generic to the following disclosed patentably distinct species: the antivirals that are disclosed but not claimed are listed on page 10 beginning at line 40 or page 13 beginning on line 46 or elsewhere in the specification. The species or groups of species are independent or distinct because these active compounds are used in different treatments, they are used to treat different specific ailments, they are used in different ways and they are used to treat different patient populations. Furthermore, they are structurally and functionally different.

Specifically, they have different functions, they have different structures and they have different effects on the body when they are administered. In addition, these species are not obvious variants of each other based on the current record.

If Applicant elects the invention of Group II or III, Applicant is further required under 35 U.S.C. 121 to elect a single disclosed species, or a single grouping of patentably indistinct species, for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable.

There is a search and/or examination burden for the patentably distinct species as set forth above because at least the following reason(s) apply:

The species are structurally and functionally different. They have different mechanisms of action and exhibit different physiological effects when they are administered.

Applicant is advised that the reply to this requirement to be complete <u>must</u> include (i) an election of a species or a grouping of patentably indistinct species to be examined even though the requirement <u>may</u> be traversed (37 CFR 1.143) and (ii) identification of the claims encompassing the elected species or grouping of patentably indistinct species, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

The election may be made with or without traverse. To preserve a right to petition, the election must be made with traverse. If the reply does not distinctly and specifically point out supposed errors in the election of species requirement, the election shall be treated as an election without traverse. Traversal must be presented at the time of election in order to be considered timely. Failure to timely traverse the requirement will result in the loss of right to petition under 37 CFR 1.144. If claims are added after

the election, applicant must indicate which of these claims are readable on the elected species or grouping of patentably indistinct species.

Should applicant traverse on the ground that the species, or groupings of patentably indistinct species from which election is required, are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing them to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the species unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other species.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which depend from or otherwise require all the limitations of an allowable generic claim as provided by 37 CFR 1.141.

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to SARAH CHICKOS whose telephone number is (571)270-3884. The examiner can normally be reached on M-F 9-6.

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If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, David Blanchard can be reached on 571-272-0827. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

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April 20, 2011

/David J Blanchard/ Supervisory Patent Examiner, Art Unit 1619